Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) Blister pack for pharmaceutical use comprising blisters containing a solid dosage form of desmopressin, or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable adjuvant, diluent or carrier, wherein said solid dosage form is adapted to prevent moisture related degradation of said desmopressin.

2. (Original) Blister pack according to claim 1, wherein said solid dosage form contains an agent that provides a pH in the range of from 3.0 to 6.2 as measured when said solid dosage form is contacted with water.

3. (Original) Blister pack according to claim 2, wherein said pH is in the range of from 3.5 to 5.5.

4. (Original) Blister pack according to claim 3, wherein said pH is in the range of from 4.0 to 5.0, preferably from 4.5 to 4.8.

5. (Currently Amended) Blister pack according to ~~any one of claims 1-4~~ claim 1, wherein said agent is an acid, preferably an acid selected from a group consisting of citric acid, hydrochloric acid and malic acid.

6. (Currently Amended) Blister pack according to ~~any one of claims 1-5~~ claim 1, wherein said blisters are composed of a material selected from PVC, PVC/PVDC blends, PE, PP, polystyrene, polyester, paper, polyamide, PET, COC, aluminium foil and blends thereof.

7. (Currently Amended) Blister pack according to ~~any one of claims 1-6~~ claim 1, wherein said solid dosage form does not comprise an enteric coating.

8. (Currently Amended) Blister pack according to ~~any one of claims 1-7~~ claim 1, wherein said solid dosage form is selected from a group consisting of tablets, granulate powder, lozenge, cachet, dry powder, capsule and wafer sheet.

9. (Original) Solid dosage form of desmopressin, or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable adjuvant, diluent or carrier, wherein said solid dosage

form comprises an agent that provides a pH in the range of from 4.5 to 5.5 as measured when said solid dosage form is contacted with water; with the proviso that said solid dosage form does not comprise fish gelatin or an enteric coating.

10. (Original) Solid dosage form according to claim 9, wherein said pH is in the range of from 4.5 to 5.0, preferably from 4.5 to 4.8.

11. (Currently Amended) Solid dosage form according to ~~any one of claims 9-10~~ claim 9, wherein said agent is an acid, preferably an acid selected from a group consisting of citric acid, hydrochloric acid and malic acid.

12. (Currently Amended) Solid dosage form according to ~~any one of claims 9-11~~ claim 9, which is selected from a group consisting of tablets, granulate powder, lozenge, cachet, dry powder, capsule and wafer sheet.

13. (Currently Amended) Blister pack for pharmaceutical use comprising blisters containing a solid dosage form as defined in ~~any one of claims 9-12~~ claim 9.

14. (Original) Blister pack according to claim 13, wherein said blisters are composed of a material selected from PVC, PVC/PVDC blends, PE, PP, polystyrene, polyester, paper, polyamide, PET, COC, aluminium foil and blends thereof.